

DNA Mixture Interpretation:
Principles and Practice in Component Deconvolution and Statistical Analysis

Interlaboratory Mixture Studies



AAFS 2008 Workshop #16
Washington, DC
February 19, 2008

John M. Butler
john.butler@nist.gov



Outline

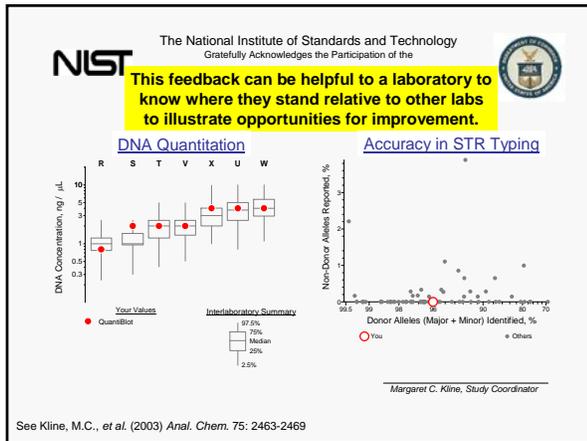
- Purpose of Interlaboratory Studies
- Overview of Mixture Studies and Lessons Learned
- NIST MIX05 Study Results

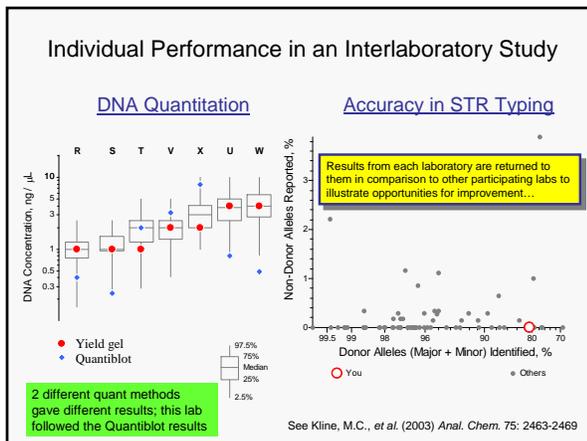
Interlaboratory Studies

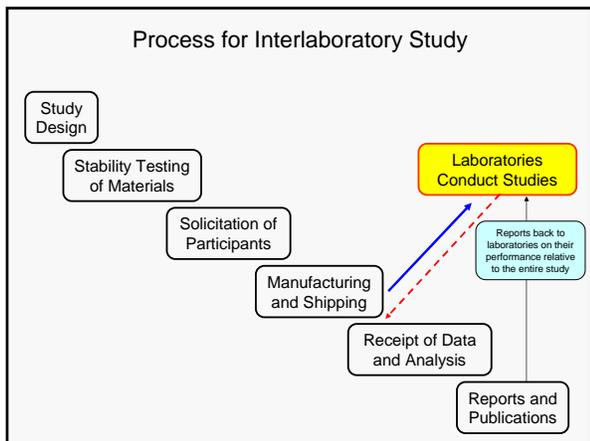
- Purpose...
 - Not a proficiency test
 - Most labs see them as opportunity to anonymously directly compare themselves to others
- STRBase section on interlab studies
 - <http://www.cstl.nist.gov/biotech/strbase/interlab.htm>

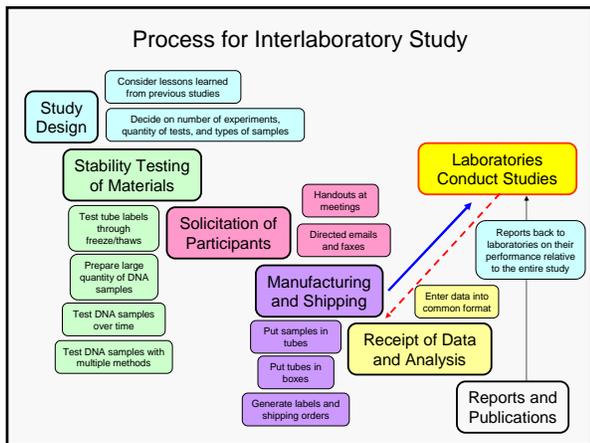
A High Degree of Variability Currently Exists with Mixture Interpretation

- “If you show 10 colleagues a mixture, you will probably end up with 10 different answers”
– Peter Gill, Human Identification E-Symposium, April 14, 2005
- Interlaboratory studies help to better understand why variability may exist between laboratories
- Most analysts are only concerned about their own lab protocols and do not get an opportunity to see the big picture from the entire community that can be provided by a well-run interlaboratory study









NIST Initiated Interlaboratory Studies		
Studies involving STRs	# Labs	Publications
Evaluation of CSF1PO, TPOX, and TH01	34	Kline MC, Duetter DL, Newall P, Redman JW, Reeder DJ, Richard M. (1997) Interlaboratory evaluation of STR triplex CTT. <i>J. Forensic Sci.</i> 42: 897-906
Mixed Stain Studies #1 and #2 (Apr–Nov 1997 and Jan–May 1999)	45	Duetter DL, Kline MC, Redman JW, Newall PJ, Reeder DJ. (2001) NIST Mixed Stain Studies #1 and #2: interlaboratory comparison of DNA quantification practice and short tandem repeat multiplex performance with multiple-source samples. <i>J. Forensic Sci.</i> 46: 1199-1210
MSS3		
Mixed Stain Study #3 (Oct 2000–May 2001)	74	Kline, M.C., Duetter, D.L., Redman, J.W., Butler, J.M. (2003) NIST mixed stain study 3: DNA quantitation accuracy and its influence on short tandem repeat multiplex signal intensity. <i>Anal. Chem.</i> 75: 2463-2469. Duetter, D.L., Kline, M.C., Redman, J.W., Butler, J.M. (2004) NIST Mixed Stain Study #3: signal intensity balance in commercial short tandem repeat multiplexes. <i>Anal. Chem.</i> 76: 6928-6934.
DNA Quantitation Study (Jan–Mar 2004) QS04	80	Kline, M.C., Duetter, D.L., Redman, J.W., Butler, J.M. (2005) Results from the NIST 2004 DNA Quantitation Study. <i>J. Forensic Sci.</i> 50(3):571-578
Mixture Interpretation Study (Jan - Aug 2005) MIX05	69	Several presentations made ... Poster at 2005 Promega meeting (Sept 2005); available on STRBase

Overall Lessons Learned
from NIST MSS 1,2,&3

- Laboratories have instruments with different sensitivities
- **Different levels of experience and training plays a part in effective mixture interpretation**
- Amount of input DNA makes a difference in the ability to detect the minor component (labs that put in “too much” DNA actually detected minor components more frequently)

**NIST MIX05
Summary**

Purpose of MIX05 Study

- **Goal is to understand the “lay of the land” regarding mixture analysis across the DNA typing community**
- One of the primary benefits we hope to gain from this study is **recommendations for a more uniform approach to mixture interpretation** and training tools to help educate the community

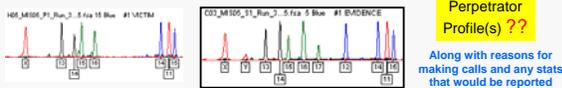
MIX05 Study Design and Purpose

Interlab studies provide a "big picture" view of the community

- Permit a large number of forensic practitioners to evaluate the same mixture data
- Provide multiple cases representing a range of mixture scenarios
- Generate data from multiple STR kits on the same mixture samples to compare performance for detecting minor components
- The primary variable should be the laboratory's interpretation guidelines rather than the DNA extraction, PCR amplification, and STR typing instrument sensitivity
- Are there best practices in the field that can be advocated to others?

Mixture Interpretation Interlab Study (MIX05)

- Only involves interpretation of data – to remove instrument detection variability and quantitation accuracy issues
- 94 labs enrolled for participation
- 69 labs have returned results (17 from outside U.S.)
- Four mock cases supplied with "victim" and "evidence" electropherograms (GeneScan .isa files – that can be converted for Mac or GeneMapper; gel files made available to FMBIO labs)
- Data available with Profiler Plus, COfiler, SGM Plus, PowerPlex 16, Identifiler, PowerPlex 16 BIO (FMBIO) kits
- Summary of results will involve training materials to illustrate various approaches to solving mixtures



Requests for Participants in MIX05

Mixtures representing four different case scenarios have been generated at NIST with multiple STR kits and provided to laboratories as electropherograms.

We would like to receive the following information:

- 1) Report the results as though they were from a real case including whether a statistical value would be attached to the results. Please summarize the perpetrator(s) alleles in each "case" as they might be presented in court—along with an appropriate statistic (if warranted by your laboratory standard operating procedure) and the source of the allele frequencies used to make the calculation. Please indicate which kit(s) were used to solve each case.
- 2) Estimate the ratio for samples present in the evidence mixture and how this estimate was determined.
- 3) Provide a copy of your laboratory mixture interpretation guidelines and a brief explanation as to why conclusions were reached in each scenario

A MIX05 Participant Noted...

“Things we do not do:

- **Calculate mixture ratios for casework**
 - **Calculation used for this study:** Find loci with 4 alleles (2 sets of sister alleles). Make sure sister alleles fall within 70%, then take the ratio of one allele from one sister set to one allele of the second sister set, figure ratios for all combinations and average. Use peak heights to calculate ratios.
- **Provide allele calls in reports**
- **Provide perpetrator(s) alleles or statistics in court without a reference sample to compare to the DNA profile obtained from the evidence. We will try to determine the perpetrator(s) profile for entry into CODIS.”**

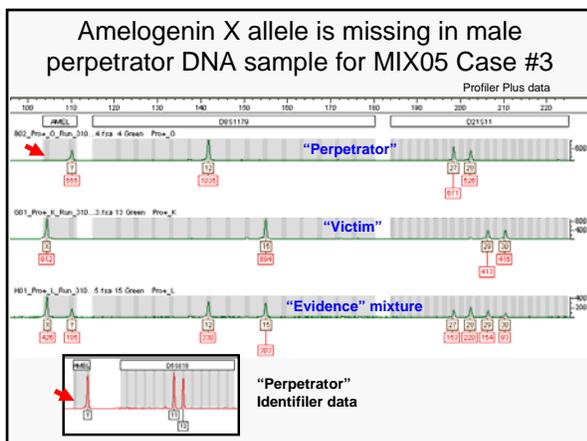
We recognize that some of the information requested in this interlab study may not be part of a lab's standard operating procedure

MIX05 Case Scenarios

Based on Identifier 15 STR loci

	#alleles		#loci with #alleles				
	N	N	N	N	N	N	N
Genomic DNA samples with specific allele combinations (“evidence”) were mixed in the following ratios:	all	unq	1	2	3	4	5
Case #1 – victim is major contributor (3F:1M)	39	26	2	6	5	2	0
Case #2 – perpetrator is major contributor (1F:3M)	55	52	0	1	4	10	0
Case #3 – balanced mixture (1F:1M) • Male lacked amelogenin X	48	37	0	3	8	4	0
Case #4 – more extreme mixture (7F:1M) • Male contained tri-allelic pattern at TPOX	50	42	0	3	7	4	1

Female victim DNA profile was supplied for each case
Labs asked to deduce the perpetrator DNA profile – suspect(s) not provided



MIX05 Results on Multiple Kits

<http://www.cstl.nist.gov/biotech/strbase/interlab/MIX05.htm>

Case 1 evidence (mixture)

Profiler Plus 

COfiler 

Identifiler 

PowerPlex 16 

SGM Plus 

ABI 3100 Generated Data was supplied on CD-ROM to labs as either .fsa files (for Gen typer NT or GeneMapperID) or Mac-converted files for Gen typer Mac

FMBIO data was also made available upon request

Summary of MIX05 Responses

94 labs enrolled for participation

69 labs returned results (17 from outside U.S.)

50 labs made allele calls

39 labs estimated ratios

29 labs provided stats

All participants were supplied with all data and could choose what kits to examine based on their experience and lab protocols

Generally Identifiler data was of poorer quality in the electropherograms we provided...which caused some labs to not return results (they indicated a desire for higher quality data through sample re-injection to reduce pull-up prior to data interpretation)

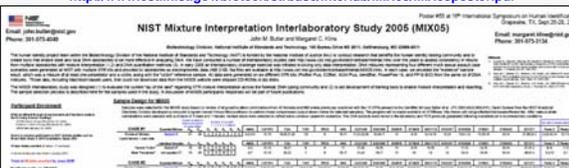
STR kit results used

- 34 ProfilerPlus/COfiler
- 10 PowerPlex 16
- 7 PP16 BIO
- 5 Identifiler
- 2 SGM Plus
- 1 All ABI kit data
- 9 Various combinations

What MIX05 Participants Have Received Back from NIST...

- Certificate of participation in the interlab study
- Copy of the poster presented at the Promega Sept 2005 meeting displaying "correct" results for the perpetrator in each case scenario as well as an explanation of study design and preliminary results

<http://www.cstl.nist.gov/biotech/strbase/interlab/MIX05/MIX05poster.pdf>



Different Stats Used

Combined Probability
of Exclusion

↑

↓

Random Match Probability
on Deduced Profiles

- **Lab 9** (4.14×10^7) used 1/CPI
- **Lab 6** (4.0×10^7) used selected loci and summed all possible genotypes for loci not completely deduced
- **Lab 90** (1.18×10^{15}) used theta value of 0.03 and deduced alleles at all 13 loci (**correctly deduced all perpetrator alleles**)

Different Thresholds of Detection Influence Allele Calls

TECHNICAL NOTE

J. Forensic Sci., January 2007, Vol. 52, No. 1
doi:10.1111/j.1744-3130.2006.00161.x
Available online at www.blackwell-synergy.com

Jason R. Gilder,¹ M.S.; Travis E. Doorn,² Ph.D.; Keith Inman,³ M. Crim.; and Dan E. Krane,⁴ Ph.D.

Run-Specific Limits of Detection and Quantitation for STR-based DNA Testing

150 RFU
LOQ (77 RFU)
LOD (29 RFU)

FIG. 3.—Electropherograms from an approximately 10:1 mixture of two reference samples. Three different thresholds are shown: a minimum peak height threshold of 150 relative fluorescence units (RFU) (dashed line); a limit of quantitation (LOQ) threshold determined to be at 77 RFU; and the negative control for this electropherogram was labeled (NC) and a limit of detection (LOD) threshold determined to be at 29 RFU. In this electropherogram, the smallest standard (i.e., Genotyper[®] assigned allele calls with 99% minor share in place) are shown in boxes immediately before the electropherogram peaks whose peak heights (in RFU) are shown to meet or exceed their labels for all peaks with heights greater than the LOD. Peaks consistent with the known profile of the minor component are marked.

Gilder, J.R., Doorn, T.E., Inman, K., Krane, D.E. (2007) Run-specific limits of detection and quantitation for STR-based DNA testing. *J. Forensic Sci.* 52(1): 97-101.

Different Detection Thresholds Used

LabID	Kits Used	Case 1 Caucasians	Notes
90	ProPlus/Cofiler	1.18E+15	75 RFUs; all 13 STRs; all results correct
34	ProPlus/Cofiler	2.40E+11	Not stated; 8 STRs, 2 partial, 3 INC
33	ProPlus/Cofiler	2.94E+08	75 RFUs; no deduced alleles reported
66	ProPlus/Cofiler	40,000,000	Not provided; 3 STRs, 6 partial, 4 INC
9	ProPlus/Cofiler	4.14E+07	100 RFUs; no deduced alleles reported
79	ProPlus/Cofiler	930,000	150 RFUs; 2 STR, 5 partial, 6 INC
16	ProPlus/Cofiler	434,600	Not stated; no deduced alleles reported

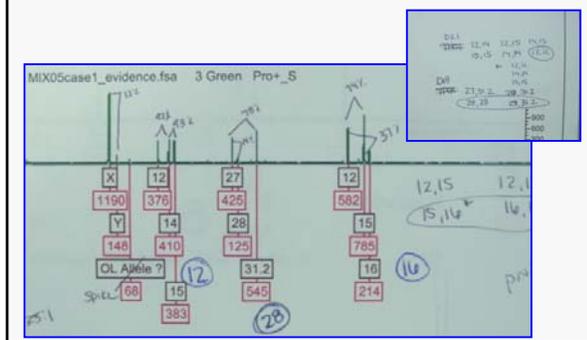
- **Lab 90** has **specific, detailed mixture interpretation guidelines** with worked examples and a fabulous flowchart
- **Lab 16** has **vague guidelines** that begin with "mixture interpretation is not always straightforward. Analysts must depend on their knowledge and experience..."

Examples of MIX05 Report Formats

All examples with Case #1

(~3:1 mixture with female victim as the major component – and victim profile is provided)

Manual Solving of MIX05 Peak Ratios and Possible Mixture Combinations



Another MIX05 Participant Manually Solving a Mixture

Q215117	13	1081	ratio = 153.95 = 284	
Q215117	14	132		24.4/150 = 0.1626
Q215111	28	972		
Q215111	30	164		
Q215111	31	88	ratio = 283	ratio = 2335 / 8.4 = 0.1132
Q215111	22.2	1010		
D185051	12	162		
D185051	10	138	ratio = 300	ratio = 2297 / 3.0 = 0.1328
D185051	17	364		
D185051	18	1033		
Q55818	8	1050		
Q55818	11	140	ratio = 372	10.6 / 28.5 = 0.372
Q55818	12	233	ratio = 328	ratio = 2328 / 7.15 = 0.1308
Q55818	13	943	ratio = 47.15	
D135317	8	129	ratio = 270	
D135317	9	141		
D135317	13	905		
D135317	14	817		ratio = 1902 / 2.1 = 0.1338
D75820	8	887	ratio = 283	ratio = 1618 / 2.1 = 0.1911
D75820	9	155		
D75820	10	805	ratio = 24	ratio = 201 / 156 = 0.129
D75820	11	98	ratio = 239	
D351358	10	1543	ratio = 218	ratio = 1857 / 8.4 = 0.1488
D351358	18	124		
D185539	9	282		
D185539	10	1420	ratio = 71	ratio = 3181 / 21.5 = 0.1278
D185539	11	1337	ratio = 404	ratio = 1337 / 10.5 = 0.1278
D185539	12	213		
T401	7	738		
T401	8	87	ratio = 168	ratio = 1087 / 6.7 = 0.1079
T401	8.5	680		
T401	10	81		

Different Reporting Formats for MIX05 Data

Profile that would be put into CODIS

LOCI	CODIS ENTRY * obligate allele	OTHER ALLELE'S IN SUSPECT'S POSSIBLE PROFILE
D3S1358	17	16,17
VWA	15*	15,17
FGA	20,22	20,22
D8S1179	12	12,12
D21S11	28*	28,31.2
D18S51	15*	15,16
D5S818	--	--
D13S317	12	12,12
D7S820	--	10
D16S539	10,11*	10,11
THO1	7*	7,8 maybe
TPOX	8	8 maybe
CSF1PO	--	11,12 maybe

Kits – Profiler Plus and Cofiler
Ratio - 1:2 (perpetrator:victim)

Different Reporting Formats for MIX05 Data

Locus	Items	
	"S" Case 1 Evid.	"P" Case 1 Victim
D3S1358	15, 16, *	15, 16
D16S539	(10), 11, (12)	11, 12
AMEL	X, *	X
THO1	(7), 8	8
TPOX	8	8
CSF1PO	11, 12	11, 12
D7S820	9, 10	9, 10
vWA	(15), 17	17
FGA	19, 20, 21, 22	19, 21
D8S1179	12, 14, 15	14, 15
D21S11	27, 31.2, *	27, 31.2
D18S51	12, 15, (16)	12, 15,
D5S818	11	11
D13S317	11, 12	11

() Indicates apparent minor peaks in a mixture.
* indicates peaks below the VFL threshold (150 rfu) for reporting.

Different Reporting Formats for MIX05 Data

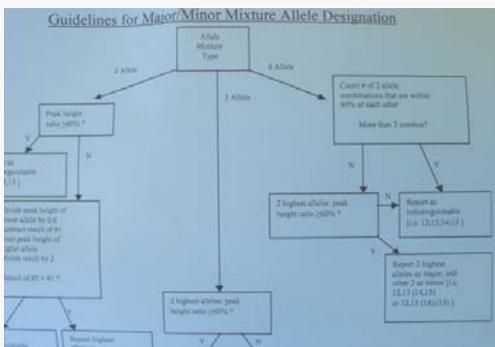
Case 1:

Item description	D3S1358	VWA	FGA	AMEL	D8S1179	D21S11	D18S51	D5S818	D13S317/D7S820	D16S539	THO1	TPOX	CSF1PO
Pro+CO S evid 1	15,16 (17)	15,17 21,22	19,20 (21)	X,X (Y)	12,14,15 (28)	27,31.2 (16)	12,15 (16)	11,11	11,12 9,10 11,12	10,11 12	7,8	8,8	11,12
Pro+CO P victim 1 reference	15,16	17,17	19,21	X,X	14,15	27,31.2	12,15	11,11	11,11 9,10 11,12	8,8	8,8	11,12	
Male interpreted from evidence 1	17	15,15 15,17	20,22	X,Y	12,12	28	16	11,11	12,12 Nd	10,11 7,7 7,8	Nd	Nd	

Two allele values separated by a comma represent a genotype. Genotype calls assume biallelic donors with no null alleles.
() Indicates minor allele detected.
Single numbers and numbers separated by "-" represent an allele only designation rather than a genotype.
Interpreted profile assumes that the victim is present in the evidence mixture of two people. More than one genotype may be listed where a single genotype could not be conclusively determined. Nd=not determined due to level of results.

The community would benefit from more uniform reporting formats and mixture solving strategies...

Some Protocols Have Flow Charts to Help Make Decisions in Mixture Resolution



Value of the MIX05 Study

<http://www.cstl.nist.gov/biotech/strbase/interlab/MIX05.htm>

- Data sets exist with multiple mixture scenarios and a variety of STR kits that **can be used for training purposes**
- A wide variety of approaches to mixture interpretation have been applied on the **same data sets evaluated as part of a single study**
- **Interpretation guidelines from many laboratories are being compared to one another for the first time in an effort to determine challenges facing future efforts to develop "expert systems" for automated mixture interpretation**
- **We are exploring the challenges of supplying a common data set to a number of forensic laboratories** (e.g., if a standard reference data set was ever desired for evaluating expert systems)

Conclusions from the MIX05 Study (Opportunities for Improvement)

- It is worth taking a closer look at protocol differences between labs to see the impact on recovering information from mixture data
- Training should help bring greater consistency
- Expert systems (when they become available and are used) should help aid consistency in evaluating mixtures and help produce more uniform reporting formats

Virtual MixtureMaker Output

	1	2	3	4	5	6	7	8
1	From	To	N ₁	N ₂	N ₃	N ₄	N ₅	N ₆
2	Caucasian WT51354	AFamer ZT79338	0	1	2	12	0	0
3	Caucasian JA16929	AFamer OT05565	0	3	3	9	0	0
4	Caucasian GT38073	AFamer MT95372	0	2	3	10	0	0
5	AFamer ZT79307	Caucasian MT97141	0	2	3	10	0	0
6	Caucasian OT07753	Hispanic GT37402	0	1	3	11	0	0
7	Hispanic GT37767	AFamer GT37019	1	7	4	3	0	0
8	AFamer ZT79330	Hispanic PT84633	0	1	4	7	0	0
9	Caucasian MT97188	AFamer OT05894	0	2	4	9	0	0
10	Caucasian MT94843	AFamer OT05568	0	1	4	10	0	0
11	AFamer ZT79338	Caucasian MT94848	0	1	4	10	0	0
12	AFamer OT05597	Hispanic TT51407	0	1	4	10	0	0

When the STR profiles for these two individuals are combined to create a 2-person mixture, the mixture profile will contain 1 locus with a single allele, 7 loci with two alleles, 4 loci with three alleles, and 3 loci with four alleles (and no loci with 5 or 6 alleles, which is only possible if one or both samples possess tri-allelic patterns at the same STR locus).

Virtual MixtureMaker Output

Female	Male	N ₁	N ₂	F ₁₀₀	F ₉₉	N ₁	N ₂	N ₃	N ₄	N ₅	N ₆	AMEL	CSFIPO	FGA	TH01	TPOX
Caucasian T150722	AFamer ZT79619	55	53	0.96	0	0	5	10	1	0	0	X,X,Y	7,10,12,13	20,23,24	7,8,9,3,10	8,9,10,11
Individual Sample	N ₁	N ₂	N ₃	N ₄	N ₅	N ₆	AMEL <td>CSFIPO <td>FGA <td>TH01 <td>TPOX <td></td> <td></td> <td></td> <td></td> <td></td> </td></td></td></td>	CSFIPO <td>FGA <td>TH01 <td>TPOX <td></td> <td></td> <td></td> <td></td> <td></td> </td></td></td>	FGA <td>TH01 <td>TPOX <td></td> <td></td> <td></td> <td></td> <td></td> </td></td>	TH01 <td>TPOX <td></td> <td></td> <td></td> <td></td> <td></td> </td>	TPOX <td></td> <td></td> <td></td> <td></td> <td></td>					
Caucasian T150722	16	31					X,X	12,13	23,24	8,10	8,11					
AFamer ZT79619	16	29					X,Y	7,10	20,24	7,9,3	9,10					
Female	Male	N ₁	N ₂	F ₁₀₀	F ₉₉	N ₁	N ₂	N ₃	N ₄	N ₅	N ₆	AMEL	CSFIPO	FGA	TH01	TPOX
Caucasian T150699	AFamer OT05588	50	45	0.90	0.87	0	3	7	4	1	1	X,X,X,Y	10,11,12,13	23,24,25	8,9,9,3	8,9,10,11,12
Individual Sample	N ₁	N ₂	N ₃	N ₄	N ₅	N ₆	AMEL <td>CSFIPO <td>FGA <td>TH01 <td>TPOX <td></td> <td></td> <td></td> <td></td> <td></td> </td></td></td></td>	CSFIPO <td>FGA <td>TH01 <td>TPOX <td></td> <td></td> <td></td> <td></td> <td></td> </td></td></td>	FGA <td>TH01 <td>TPOX <td></td> <td></td> <td></td> <td></td> <td></td> </td></td>	TH01 <td>TPOX <td></td> <td></td> <td></td> <td></td> <td></td> </td>	TPOX <td></td> <td></td> <td></td> <td></td> <td></td>					
Caucasian T150699	15	27					X,X	10,12	23,24	9,3	8,12					
AFamer OT05588	15	27					X,Y	11,13	26	8,9	8,10,11					

→ One locus with 5 alleles in this 2-person mixture
→ 16 loci examined with 31 distinguishable alleles
→ No locus failures in this profile
→ One tri-allelic locus
→ 13 heterozygous loci
→ 2 homozygous loci

Some Final Thoughts...

- It is of the highest importance in the art of detection to be able to recognize out of a number of facts, which are incidental and which vital. Otherwise your energy and attention must be dissipated instead of being concentrated (Sherlock Holmes, *The Reigate Puzzle*).
- **“Don’t do mixture interpretation unless you have to”** (Peter Gill, Forensic Science Service, 1998).
- Mixture interpretation consumes a large part of DNA analysts’ time – software tools that improve consistency in analysis will speed casework reporting and hopefully cases solved



Forensic Science International 134 (2003) 130–136

www.elsevier.com/locate/forensic



DNA mixtures in forensic casework: a 4-year retrospective study

Yolanda Torres^{a,*}, Inmaculada Flores^a, Victoria Prieto^a, Manuel López-Soto^a,
 María José Farfán^a, Angel Carracedo^a, Pilar Sainz^b

^aInstituto Nacional de Toxicología y Forensic, A. Postal 961, E-41010 Sevilla, Spain
^bInstituto de Medicina Legal, Universidad de Santiago de Compostela, Sevilla, Spain

Received 27 March 2003; accepted 1 April 2003

Conclusion

"Mixture interpretation theory is well established and used in forensic laboratories. Most mixtures detected in casework are satisfactorily solved. But from this revision we can conclude that the behaviour of each mixed sample can be different and multifactorial and occasionally its interpretation turns out to be complicated—sometimes paralleling the importance of the evidence in the resolution of the case. In some casework mixtures our experience has proved that theoretical assumptions from studies with laboratory samples, albeit very useful, can turn out to be impracticable. We consider that more sharing of day to day forensic laboratory problems is needed to refine our technical procedures in the resolution of specially difficult evidence."

Acknowledgments

Funding from interagency agreement 2003-IJ-R-029 between NIJ and the NIST Office of Law Enforcement Standards

NIST Human Identity Project Team – Leading the Way in Forensic DNA...









John
Butler

Margaret
Kline

Pete
Vallone

Jan
Redman

Amy
Decker

Becky
Hill

Dave
Duewer

Role in MIX05

- Margaret Kline (running study, sample prep, data review)
- John Butler (study design and data review)
- Becky Hill (GeneMapperID data review)
- Jan Redman (Access database entry, shipping)
- Dave Duewer (*Virtual MixtureMaker* to aid sample selection; **mixSTR program**)
- Chris Tomsey & Frank Krist (FMBIO Mac data)
- Kermit Channel & Mary Robnett (FMBIO NT data)

The many forensic scientists and their supervisors who took time out of their busy schedules to examine the MIX05 data provided as part of this interlaboratory study
